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(54) Automatic injection devices

Automatische Injektionsvorrichtung Dispositif d'injection automatique

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(56) References cited:

DE-A- 4 037 418 GB-A- 728 248

FR-A- 1 538 565

US-A-4 998 922

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and thus made safe after use.

This invention relates to injection devices, and in particular medical ones where the needle is retracted

Such injection devices are in increasing demand for obvious reasons. There is great danger in a discarded syringe with a possibly infected needle. It is highly desirable that retraction of the needle after use should not only be possible but also be automatic. It should not rely on the user making it safe by some manipulation which can all too easily be overlooked. Also, with the increasing use of self-administered drug therapy, there is a demand for a device whereby the syringe needle is not normally visible to the user before or after the injection.

In GB-A-728248 hypodermic injection apparatus is described which meets these requirements. But the various embodiments are of considerable complexity, and therefore expense, and it is believed that none of them ever came into common use. Even though the apparatus was safe before and after injection, that is now not enough: it should preferably be cheap enough to be used once and then disposed of.

The apparatus of GB-A-728248 employs a powerful spring which, when released, thrusts forward on an ampoule and needle carrier. This projects the needle into the flesh of the patient. At the end of this phase, the spring is automatically decoupled from the carrier but continues to act on a plunger which co-operates with a piston within the ampoule. Thus the contents of the ampoule are squeezed out through the needle by the spring. At the end of this phase, the spring is automatically decoupled from the plunger, leaving the ampoule and needle carrier free to be acted upon by a relatively weak return spring, which urges the carrier back to a retracted position.

There are therefore two decouplings and the complexity referred to above is largely attributable to these. It is the aim of this invention to eliminate one of these decoupling arrangements and thereby substantially simplify the construction.

According to the present invention there is provided an injection device comprising a barrel, a spring loaded drive member therein, release mechanism for allowing said member to be sprung forwardly within the barrel, a spring loaded, charged capsule within the barrel, a needle associated with the capsule, initially in a retracted position at the forward end of the barrel, the spring loading of the capsule being in opposition to but weaker than the spring loading of said drive member, a plunger with which the forward end of said drive member co-operates, and means for decoupling the plunger from said drive member at the end of the forward stroke, the arrangement being such that said forward stroke first drives the capsule and the needle to a needle projecting position, and secondly forces the plunger to eject the capsule charge through the needle, the capsule spring loading being then freed by the decoupling means to return the capsule and needle to the needle retracted position,

characterised in that for the first part of the forward stroke said drive member acts through the plunger and the capsule charge.

Thus, no decoupling is required at the end of the first phase; the spring loaded drive member having relatively easily projected the needle carries on more slowly against the resistance of the capsule charge whose only escape is through the needle.

The capsule will conveniently be a proprietary syringe with its plunger removed and replaced by said plunger which co-operates with said drive member.

Preferably, the spring loading will be provided by coil springs co-axial within the barrel.

In one form, the decoupling means is provided by the rear end of said plunger which is abutted by said drive member and which is adapted to be deformed when it comes into co-operation with the capsule. For example, the rear end of the plunger may be bifurcated and there may be exterior projections on the fingers formed thereby, said projection when engaged by the capsule on entry therein causing the fingers to be wedged together and in this constricted condition to be free to enter a passage within the drive member.

In an alternative arrangement, the decoupling means may be provided by a thrust device between the rear end of the plunger and the forward end of the drive member, and a contoured passage within the barrel, the thrust device being maintained effectively rigid by confinement within a narrow portion of said passage for all but the final stage of the forward stroke, but being rendered non-rigid when the confinement is eased by a wide portion of said passage for said final stage. One example of such a thrust device is an element such as a ball that rolls free or otherwise escapes from between the plunger and the drive member when it attains the freedom of said wide portion. Another possible thrust device is a hinged linkage which is maintained substantially straight and extended by said narrow portion but which collapses sideways when it attains the freedom of said wide por-

The drive member will generally have a catch which initially is engaged with the barrel, holding the forward spring energised. The release mechanism, which may be a button-like rear end cap, frees this catch.

For a better understanding of the invention, some embodiments will now be described, by way of example, with reference to the accompanying drawings, in which:

Figures 1-4 are longitudinal sections of an injection device in progressive stages of operation,

Figure 5 is a sectional detail of an alternative injection device, and

Figure 6 is a sectional detail of a third injection device.

The injection device of Figures 1 to 4 has a stepped barrel 1 reducing towards the forward end where there is a cap 2 which is removed for use. At the rear end there is a cap 3 which is snapped on by its peripheral flange

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and which has an inner annulus 4 and a central tab 5. The cap 3 is rotatable on the barrel 1 and its central portion can be flexed inwards, axially of the barrel, by virtue of an annular weakness 6 near the periphery.

The largest diameter portion 7 of the barrel houses a bottle shaped drive member 8 whose neck is towards the rear and which has a central aperture in its base. The neck 9 is bifurcated and extends through an aperture 10 defined by an inwardly extending flange 11. Beyond that flange the drive member 8 terminates in out-turned hooks with bevelled surfaces 12 with which the annulus 4 cooperates. At the other, forward end of the drive member 8, there is an outward flange 13, and a coil spring 14 surrounding the member acts between this and the flange 11. As shown in Figure 1, with the drive member 8 captive to the rear of the barrel, this spring is at its maximum compression.

The transition between the rear portion 7 and the intermediate portion 15 of the barrel 1 is internally defined by an annular rib 16. At the forward end of this portion 15 there is another annular rib 17 and initially a charged capsule 18 is located by these ribs, a surrounding coil spring 19 reacting against the rib 17 and urging a flange 20 at the rear of the capsule against the rib 16. The spring 19 is somewhat weaker than the spring 14 and the forward travel of the capsule, when this spring 19 becomes compressed, is limited by spacers 21 extending rearwardly from the rib 17 to provide an abutment for the flange 20.

The leading portion of the capsule 18, with a needle 22, is located in the forward end portion 23 of the barrel, the tip of the needle being set back from the end. Initially, the needle 22 is sheathed in a self-sealing silicon rubber shroud 24, protecting against contamination and leakage. The shroud is captive within an inward tubular extension 25 of the cap 2 and so is pulled off when that cap is removed just prior to use.

A plunger 26 is formed by a rod 27 and a piston 28 within the capsule 18. They are not connected: the piston is provided with the capsule 18, which may be filled with its charge and plugged by the piston quite separately from the assembly of the syringe. The rear end of the rod 27 is bifurcated and near the extreme end the resultant fingers are enlarged to form exterior shouldered abutments 29 against which the annular base of the drive member 8 acts. Near the root of the bifurcation there are further enlargements 30 on opposite sides with sloping surfaces which will wedge into the rear end of the capsule 18, as described below, to close the tips of the fingers together.

For use, the cap 2 is removed taking with it the shroud 24, and the cap 3 is turned through 90° from the "safe" position in Figure 1, where the tab 5 is holding the hooks 12 apart and firmly engaged with the flange 11. The cap 3 is shown in the "use" position in Figures 2, 3 and 4. The device is then applied to where the injection is to be made and the centre of the cap 3 is pressed. The annulus 4 wedges the hooks 12 together and frees them from the flange 11. The spring 14 is then free to act and

it shoots the drive member 8 forwards. The fluid in the capsule is virtually incompressible and it has a very narrow means of escape through the needle 22. The plunger 26 is therefore acting on a substantially solid body and it carries the capsule 18 forwards, compressing the spring 19. This action terminates as shown in Figure 2, with the needle 18 projecting and the flange 20 hard up against the spacers 21.

With the capsule 18 arrested, the plunger 26 carries on under the influence of the dominant spring 14, forcing liquid out through the needle 22. As it reaches the end of the forward stroke (Figure 3) the wedges 30 act to squeeze the bifurcated end of the rod 27 together, thus bringing the abutments 29 within the compass of the aperture at the base of the drive member 8. At this point, the drive member 8 is arrested by the rib 16, and so the spring 19 can then act, carrying the capsule 18 back with the rod 27 passing into the drive member 8 until the flange 20 abuts the forward side of the rib 16 (Figure 4). This is the initial position of the capsule, and the needle is safe within the barrel.

Referring now to Figure 5, a spring loaded drive member 31 acts on a plunger rod 32 through the intermediary of a toggle linkage 33. The toggles 34 and end discs 35 and 36 may be integrally moulded of plastics material with weaknesses at the hinge points. Initially, the toggles 34 are confined in a passage 37 so that they are virtually straight, as shown in full lines. However, towards the end of the forward stoke, the rear end of the rod 32 reaches an enlarged passage 38, and when the linkage 33 follows it is free to expand sideways as shown in broken lines. Thus, the thrust on the rod 32 is finished and the return spring acts on the plunger/capsule assembly. The rod 32 is urged rearwardly closing the two discs 35 and 36 together, this travel being sufficient to withdraw the needle.

Referring now to Figure 6, a similar configuration of passages is used, with a narrow one 40 widening into an enlarged one 41. Adrive member 42 has a cone 43 pointing forwards, its vertex engaging a ball 44 which is just slightly less in diameter than the passage 40. A similar cone 45 is at the rear end of a plunger rod 46, its vertex pointing to the rear also to engage the ball 44.

On the forward stroke the ball is restricted by the passage 40, and so transmits the thrust from the drive member 42 to the plunger rod 46. However, at the end of the forward stroke the coned end 45 enters the enlarged passage 41 and when it reaches the broken line position, the ball 44, which is in unstable captivity, falls away to one side, also as shown in broken lines. Thus the plunger, with the capsule, is free to return.

In order to facilitate construction and assembly, the barrel 1 may be made in two parts joined together after insertion of the capsule 18. The junction may conveniently be in the region of the rib 16.

In Figures 5 and 6, in order to achieve sufficient needle travel, the devices may have to be larger than that of Figures 1 to 4. But this can be alleviated to some extent by making those devices of flattened oval cross-section, the cross sections of the Figures being in the planes of the major axes.

Drugs are currently available pre-packaged in cartridges, similar to conventional syringes but with no end flanges. Such a cartridge has a cap with a pierceable 5 rubber membrane at the reduced diameter delivery end which accepts a double-ended needle. This is packaged inside a shield which enables it to be fitted safely and easily, as well as giving it temporary protection. To provide a flange that would make it usable with the device described above, the cartridge may be fitted inside a sleeve-like carrier. This would be similarly contoured, with a neck surrounding the delivery end of the cartridge and an outwardly projecting flange performing the function of the flange 20 at the opposite end. The neck could be threaded to retain the needle.

Claims

- 1. An injection device comprising a barrel (1), a spring 20 loaded drive member (8;31;42) therein, release mechanism (3,4,12) for allowing said member to be sprung forwardly within the barrel (1), a spring loaded, charged capsule (18) within the barrel (1), a needle (22) associated with the capsule (18) initially in a retracted position at the forward end of the barrel (1), the spring loading (19) of the capsule being in opposition to but weaker than the spring loading (14) of said drive member (8;31;42), a plunger (26;32;46) with which the forward end of said drive member (8;31;42) co-operates, and means (30;33;40) for decoupling the plunger (26;32;46) from said drive member (8;31;42) at the end of the forward stroke, the arrangement being such that said forward stroke first drives the capsule (18) and the needle (22) to a needle projecting position, and secondly forces the plunger (26;32;46) to eject the capsule charge through the needle (12), the capsule spring loading (19) being then freed by the decoupling means (30;33;40) to return the capsule (18) and needle (22) to the needle retracted position, characterised in that for the first part of the forward stroke, said drive member (8;31;42) acts through the plunger (26;32;46) and the capsule charge.
- 2. A device as claimed in Claim 1, characterised in that the capsule (18) is a proprietary syringe with its plunger removed and replaced by said plunger (25;32;46) which co-operates with said drive member (18).
- 3. A device as claimed in Claim 1 or 2, characterised in that the spring loading (14,19) is provided by coil springs co-axial within the barrel.
- 4. A device as claimed in Claim 1, 2 or 3, characterised in that the decoupling means (30) is provided by the rear end of said plunger (26) which is abutted by said drive member and which is adapted to be deformed

when it comes into co-operation with the capsule (18).

- 5. A device as claimed in Claim 4, characterised in that said rear end of the plunger (26) is bifurcated and there are exterior projections (30) on the fingers formed thereby, said projections (30), when engaged by the capsule (18) on entry therein, causing the fingers to be wedged together and in the constricted condition to be free to enter a passage within the drive member (14).
- 6. A device as claimed in Claim 1, 2 or 3, characterised in that the decoupling means is provided by a thrust device (33;44) between the rear end of the plunger (32:46) and the forward end of the drive member (31;44), and a contoured passage (37;38;40;41) within the barrel, the thrust device (33:44) being maintained effectively rigid by confinement within a narrow portion (37;40) of said passage for all but the final stage of the forward stroke, but being non-rigid when the confinement is eased by a wide portion (38;41) of said passage for said final stage.
- 7. A device as claimed in Claim 6, characterised in that said thrust device is an element (44) that rolls free or otherwise escapes from between the plunger (46) and the drive member (42) when it attains the freedom of said wide portion (41).
- 8. A device as claimed in Claim 6, characterized in that said thrust device is a hinged linkage (33) which is maintained substantially straight and extended by said narrow portion but which collapses sideways when it attains the freedom of said wide portion (38).

Patentansprüche

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1. Injektionsgerät mit einem Zylinder (1), einem darin angeordneten federbelasteten Antriebselement (8;31;42), einem Auslösemechanismus (3,4,12), um dem Antriebselement ein Vorschnellen im Zylinder (1) zu gestatten, einer federbelasteten, gefüllten Kapsel (18) innerhalb des Zylinders (1), wobei die auf die Kapsel wirkende Feder (19) entgegengesetzt zu der auf das Antriebselement (8;31;42) wirkenden Feder (14) wirkt und schwächer als diese ist, einer an die Kapsel (18) angeschlossenen Nadel (22), die sich anfänglich in einer zurückgezogenen Stellung am vorderen Ende des Zylinders (1) befindet, einem Stößel (26;32;46), mit dem das vordere Ende des Antriebselements (8:31:42) zusammenwirkt, und einer Vorrichtung (30;33;40) zum Entkopdes Stößels (26;32;46) von Antriebselement (8;31;42) am Ende des Vorwärtshubes, wobei die Anordnung derart ist, daß beim Vorwärtshub zuerst die Kapsel (18) mit der Nadel (22) in eine vorstehende Position gebracht wird und dann der Stößel (26;32;46) den Inhalt der Kapsel 10

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durch die Nadel (22) ausschiebt, worauf die auf die Kapsel (18) wirkende Feder (19) von der Entkopplungsvorrichtung (30;33;40) freigegeben wird, um die Kapsel (18) und die Nadel (22) in die zurückgezogene Stellung zurückzuführen, dadurch gekenn- 5 zeichnet, daß für den ersten Teil des Vorwärtshubes das Antriebselement (8;31;42) über den Stößel (26:32:46) und den Inhalt der Kapsel (18) wirksam ist.

- 2. Injektionsgerät nach Anspruch 1, dadurch gekennzeichnet, daß die Kapsel (18) eine handelsübliche Spritze ist, deren Kolben durch den mit dem Antriebselement (8;31;42) zusammenwirkenden Stößel (26;32;46) ersetzt ist.
- 3. Injektionsgerät nach Anspruch 1 oder 2, dadurch gekennzeichnet, daß die Federn (14,19) koaxial im Zylinder (1) angeordnete Schraubenfedern sind.
- 4. Injektionsgerät nach Anspruch 1, 2 oder 3, dadurch gekennzeichnet, daß die Entkoppelungsvorrichtung (30) von dem rückwärtigen Ende des Stößels (26) gebildet ist, an dem das Antriebselement anliegt und das verformt wird, wenn es mit der Kapsel (18) zsammenwirkt.
- 5. Injektionsgerät nach Anspruch 4, dadurch gekennzeichnet, daß das rückwärtige Ende des Stößels (26) gegabelt ist und daß an den dadurch gebildeten 30 Fingern äußere Vorsprünge (30) vorgesehen sind, die, wenn sie beim Eintritt in die Kapsel (18) mit dieser zusammenwirken, die Finger zusammenklemmen, so daß diese in eine Öffnung im Antriebselement (8) eintreten können.
- 6. Injektionsgerät nach Anspruch 1, 2 oder 3, dadurch gekennzeichnet, daß die Entkopplungsvorrichtung von einer Druckübertragungsvorrichtung (33;44) zwischen dem rückwärtigen Ende des Stößels (32;46) und dem vorderen Ende des Antriebselements (31;44) und von einem konturierten Kanal (37,38; 40,41) gebildet ist, wobei die Druckübertragungsvorrichtung (33;44) durch Begrenzung in einem engen Abschnitt (37;40) des Kanals für alle außer der letzten Stufe des Vorwärtshubes effektiv starr gehalten ist, jedoch für die letzte Stufe unstarr wird, wenn die Begrenzung durch einen weiten Abschnitt (38;41) des Kanals gelockert wird.
- 7. Injektionsgerät nach Anspruch 6, dadurch gekennzeichnet, daß die Druckübertragungsvorrichtung ein Element (44) ist, das zwischen dem Stößel (46) und dem Antriebselement (42) frei rollt, jedoch zwischen diesen Teilen austritt, wenn es in den Bereich des 55 weiten Abschnittes (41) kommt.
- 8. Injektionsgerät nach Anspruch 6, dadurch gekennzeichnet, daß die Druckübertragungsvorrichtung ein

Knickhebelgestänge (33) ist, daß durch den engen Abschnitt (37) im wesentlichen gerade und gestreckt gehalten wird, jedoch seitwärts zusammenklappt, wenn es in den Bereich des weiten Abschnittes (38) kommt.

Revendications

- Dispositif d'injection comprenant un cylindre (1), un élément d'entraînement (8; 31; 42) chargé par ressort dans celui-ci, un mécanisme de libération (3, 4, 12) pour permettre audit élément d'être projeté vers l'avant à l'intérieur du cylindre (1), une capsule (18) ayant une charge, chargée par ressort, à l'intérieur du cylindre, une aiguille (22) associée à la capsule (18), initialement dans une position rétractée à l'extrémité avant du cylindre (1), la charge par ressort (19) de la capsule étant en opposition à, mais plus faible que, la charge par ressort (14) dudit élément d'entraînement (8; 31; 42), un piston de compression (26; 32; 46) avec lequel coopère l'extrémité avant dudit élément d'entraînement (8; 31; 42), et un moyen (30; 33; 40) pour découpler le piston de compression (26; 32; 46) dudit élément d'entraînement (8; 31; 42) à la fin de la course vers l'avant, la disposition étant telle que ladite course vers l'avant entraîne tout d'abord la capsule (18) et l'aiguille (22) dans une position de projection de l'aiquille, et ensuite oblige le piston de compression (26; 32; 46) à éjecter la charge de la capsule à travers l'aiguille (22), la charge de ressort (19) de la capsule étant ensuite libérée par le moyen de découplage (30; 33; 40) pour ramener la capsule (18) et l'aiguille (22) à la position rétractée de l'aiguille, caractérisé par le fait que, pendant la première partie de la course vers l'avant, ledit élément d'entraînement (8; 31; 42) agit par l'intermédiaire du piston de compression (26; 32; 46) et de la charge de la capsule.
- 2. Dispositif selon la revendication 1, caractérisé par le 40 fait que la capsule (18) est une seringue de marque avec son piston de compression retiré et remplacé par ledit piston de compression (25; 32; 46) qui coopère avec ledit élément d'entraînement (8).
 - Dispositif selon la revendication 1 ou 2, caractérisé par le fait que la charge par ressort (14, 19) est fournie par des ressorts hélicoïdaux coaxiaux à l'intérieur du cylindre.
 - Dispositif selon l'une des revendications 1, 2 ou 3. caractérisé par le fait que le moyen de découplage (30) est fourni par l'extrémité arrière dudit piston de compression (26) contre lequel ledit élément d'entraînement vient en butée et qui est adapté pour se déformer lorsqu'il vient en coopération avec la capsule (18).

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- 5. Dispositif selon la revendication 4, caractérisé par le fait que ladite extrémité arrière du piston de compression (26) comporte deux branches et qu'il existe des saillies extérieures (30) sur les doigts formés de cette façon, lesdites saillies (30), lorsqu'elles sont 5 engagées par la capsule (18) à l'entrée dans celleci, amenant les doigts à se resserrer et, dans l'état resserré, à être libres d'entrer dans un passage à l'intérieur de l'élément d'entraînement (8).
- 6. Dispositif selon l'une des revendications 1, 2 ou 3, caractérisé par le fait que le moyen de découplage est fourni par un dispositif de poussée (33; 44) entre l'extrémité arrière du piston de compression (32; 46) et l'extrémité avant de l'élément d'entraînement (31; 15 42), et un passage profilé (37; 38; 40; 41) à l'intérieur du cylindre, le dispositif de poussée (33; 44) étant maintenu effectivement rigide par confinement à l'intérieur d'une partie étroite (37; 40) dudit passage pendant la totalité excepté le stade final de la course 20 vers l'avant, mais étant rendu non-rigide lorsque le confinement est relâché par une partie large (38; 41) dudit passage pendant ledit stade final.
- 7. Dispositif selon la revendication 6, caractérisé par le 25 fait que ledit dispositif de poussée est un élément (44) qui roule librement ou autrement s'échappe d'entre le piston de compression (46) et l'élément d'entraînement (42) lorsqu'il atteint la liberté de ladite partie large (41).
- 8. Dispositif selon la revendication 6, caractérisé par le fait que ledit dispositif de poussée est un organe de liaison articulé (33) qui est maintenu sensiblement rectiligne et étendu par ladite partie étroite, mais qui 35 s'affaisse latéralement lorsqu'il atteint la liberté de ladite partie large (38).

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